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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/582,459	05/16/2007	Paul Wallace	14113-00040-US	1128
	7590 06/22/201 SOVE LODGE & HUT	EXAM	IINER	
P O BOX 2207			CLARK, GREGORY D	
WILMINGTON, DE 19899			ART UNIT	PAPER NUMBER
			1786	
			MAIL DATE	DELIVERY MODE
			06/22/2010	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)		
	10/582,459	WALLACE ET AL.		
Office Action Summary	Examiner	Art Unit		
	GREGORY CLARK	1786		
The MAILING DATE of this communi Period for Reply	cation appears on the cover sheet wi	th the correspondence address		
A SHORTENED STATUTORY PERIOD FOWHICHEVER IS LONGER, FROM THE M. - Extensions of time may be available under the provisions after SIX (6) MONTHS from the mailing date of this comm - If NO period for reply is specified above, the maximum sta - Failure to reply within the set or extended period for reply Any reply received by the Office later than three months at earned patent term adjustment. See 37 CFR 1.704(b).	AILING DATE OF THIS COMMUNION of 37 CFR 1.136(a). In no event, however, may a runication. tutory period will apply and will expire SIX (6) MON will, by statute, cause the application to become AE	CATION. Peply be timely filed THS from the mailing date of this communication. ANDONED (35 U.S.C. § 133).		
Status				
1) Responsive to communication(s) file	tb) This action is non-final. for allowance except for formal matt	· •		
Disposition of Claims				
4) ☐ Claim(s) 1-27 is/are pending in the a 4a) Of the above claim(s) 6,8,11 and 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) 1-5, 7, 9-10 and 13-27 is/are 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) are subject to restrict	12 is/are withdrawn from considerare rejected.	ion.		
Application Papers				
9) The specification is objected to by the 10) The drawing(s) filed on is/are: Applicant may not request that any object Replacement drawing sheet(s) including 11) The oath or declaration is objected to	a) accepted or b) objected to attention to the drawing(s) be held in abeyare the correction is required if the drawing	ce. See 37 CFR 1.85(a). (s) is objected to. See 37 CFR 1.121(d).		
Priority under 35 U.S.C. § 119				
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 				
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (P' 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	TO-948) Paper No(s	ummary (PTO-413))/Mail Date Iformal Patent Application 		

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DETAILED ACTION

The examiner acknowledges the receipt of applicants' arguments/ amendments dated 05/20/2010. Claims 1-5, 7, 9-10, 13-27 pending.

Rejections and objections made in previous office action that do not appear below have been overcome by applicant's amendments and therefore the arguments pertaining to these rejections/objections will not be addressed.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

- (a) the invention was known or used by others in this country, or patented or described in a printed publication in this or a foreign country, before the invention thereof by the applicant for a patent.
- 1. Claims 1, 4-5, 7, 9-10, 13-18, 20, 24 and 27 are rejected under 35 U.S.C. 102(b) as being anticipated by Towns (WO 03/035714).
- 2. **Regarding Claims 1, 24 and 27,** the applicant claims an optionally substituted oligomer of polymer comprising a repeat unit of Formula 1:

$$\left(\begin{array}{cccc}
Ar^{1} - A - Ar^{2} - Ar^{2} - Ar^{3} \\
Ar^{3} & Ar^{3}
\end{array}\right)$$

Formula 1

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Wherein n is at least 1; each A is N or P; each Ar1 and Ar3 is arylene or heteroarylene; Ar2 is arylene or heteroarylene containing a linking ring to which the two A atoms are both directly linked and at least one of Ar1 or Ar2 is substituted with at least one substituent. Applicant further claims a second repeating unit wherein the second repeat unit is selected from optionally substituted phenyl, fluorene, spirobifluorene, indenofluorene, heteroaryl or dihydrophenanthrene.

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Towns discloses formula T-1 (page 4):

$$B_{1} - \begin{pmatrix} A_{1} \\ A_{2} \end{pmatrix} - \begin{pmatrix} A_{1} \\ A_{2}$$

Formula T-1 shows n =1, N corresponds to applicants' A group, Ar1 corresponds to applicants' Ar1, Ar2 corresponds to applicants' Ar2 and Ar3 corresponds to applicants' Ar3. The Ar2 linking group connects both N atoms. Ar3 can be substituted with an alkyl group or perfluoroalkyl group (page 4).

Formula T-1 also meets the limitations of claim 24.

Towns also discloses that Formula T-1 can also be a part of the copolymer T-2 (page 11):

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Copolymer T-2 shows the second repeat unit is a fluorenyl monomer (per claims 1 and 27).

- 3. **Regarding Claims 4 and 5,** Formula T-1 shows Ar1-Ar3 as phenyl groups.
- 4. **Regarding Claims 7 and 9,** copolymer T-2 shows the monomer 1 conjugated to monomer 2 (per claim 7) and Ar3 can be substituted with an alkyl group or perfluoroalkyl group (page 4) (per claim 9).
- 5. **Regarding Claim 10,** Towns discloses a blend of copolymer T-2 and another organic compound (page 13). Copolymer T-2 can be used as hole transporting material (page 2).
- 6. **Regarding Claims 13 and14,** Towns show that Formula T-1 (shown above) can have bromide as a leaving group (LG) (page 4). Copolymer T-2 (above) shows the second repeat unit is a fluorenyl monomer. Formula T-1 shows n =1, N corresponds to applicants' A group, Ar1 corresponds to applicants' Ar1, Ar2 corresponds to applicants' Ar2 and Ar3 corresponds to applicants' Ar3. The Ar2 linking group connects both N atoms. Ar3 can be substituted with an alkyl group or perfluoroalkyl group (page 4).

The reactions can be catalyzed by palladium (variable oxidation state metal) (page 6) (per claim14).

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7. **Regarding Claim 15,** Towns discloses that reactions can be catalyzed by palladium (variable oxidation state metal) (page 6) in the presence of a base (page 8).

- 8. **Regarding Claim 16,** Towns discloses that in a monomer one LG can be a reactive boron group and the other LG can be a halogen (page 8).
- 9. **Regarding Claims 17-18 and 20,** Towns discloses that the polymers or blend (page 13) (per claim 18) can be used in an optical device which can be an electroluminescent device (page 12) (per claims 17 and 20). The polymers or blend is located between two electrodes (page 13) (per claim19).

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

- (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 10. Claims 2-3 and 25-26 are rejected under 35 U.S.C. 103(a) as being unpatentable over Towns (WO 03/035714).

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11. **Regarding Claims 2-3 and 25-26** Town discloses that Ar3 in Formula T-1 (above) can be substituted with an alkyl group (page 3) or perfluoroalkyl (fluoroalkyl) (page 4). Town fails to mention substitution at Ar1 and Ar2.

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Formula T-1 is viewed as a structural isomer to the claimed compounds with substitution at Ar3. Substitution at Ar1, Ar2 or Ar3 would be bonded to aromatic rings that are in conjugation with each other. As a result, the substituents disclosed by Towns at Ar3 would be expected to give similar properties and function in a similar capacity to monomers with substitution at Ar1 or Ar2.

In the absence of unexpected result, at the time of the invention, it would have been obvious for a person of ordinary skill in the art to have made a series of monomers with varying levels of substitution at Ar1, Ar2 and Ar3 which would have included substitution at Ar1 or Ar2 with a reasonable expectation of success.

- 12. Claims 21 and 23 are rejected under 35 U.S.C. 103(a) as being unpatentable over Towns (WO 03/035714) in view of Allen (US 6,858,703).
- 13. **Regarding Claim 21 and 23,** Towns discloses that the copolymers can be used in an electroluminescent device (pages 12 and 13) but fails to mention a switching device.

Allen discloses that triarylamine polymers are used in optical sensor, switching devices and field effect transistors (column 83, lines 20-30).

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It would have been obvious to a person of ordinary skill in the art at the time of the invention to have used the electroluminescent device of Towns in applications taught by the prior art which would have included the applications disclosed by Allen which reads on applicants' claimed applications.

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- 14. Claim 22 is rejected under 35 U.S.C. 103(a) as being unpatentable over Towns (WO 03/035714) in view of Allen (US 6,858,703) and further in view of Hirai (US 6,740,900).
- 15. **Regarding Claim 22,** Towns in view of Allen disclose triarylamine polymers used as a field transistors but fail to mention the structure of the device.

Hirai discloses that an insulator layer can be placed in various locations with respect to the gate, drain and source electrodes in an organic thin-film transistor.

Hirai discloses that the organic thin-film transistor contains a organic semiconductor layer 3, a gate electrode G, a drain electrode D, and a source electrode S that are in Figure I(a) shown below:



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Hirai further discloses that a dielectric layer (field-effect transistor) serves as gate insulation layer 2 (column 7, lines 44-45). The above figure shows that the gate electrode G and the organic semiconductor layer 3 are both in contract with the gate insulation layer 2 located on side one. In addition, the above figure shows that the source electrode S and the drain electrode D are located on the organic semiconductor layer 3 on side two.

Hirai shows in the prior art that the claimed structure can be use as a field effect transistor.

With the reasonable expectation of success a person with ordinary skill in the art at the time of the invention would have selected from known field-effect transistor structures and readily substituted the conductive polymers disclosed by Towns (T-2) for the organic semiconductive layer 3 disclosed by Hirai in Figure I (a) since the conductive polymers of Towns would be functional equivalents to the polymers typically used to make an organic semiconductive layer.

- 16. Claims 1-5, 7, 9-10, 13-20 and 24-27 are rejected under 35 U.S.C. 103(a) as being unpatentable over Antoniadis (US 5,948,552) in view of Bradley (Advanced Materials, 1999, Vol. 11, No. 3, pages 241-246).
- 17. **Regarding Claims 1, 7, 24 and 27**, the applicant claims an optionally substituted oligomer or polymer containing a repeat unit of Formula 1:

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$$\left(\begin{array}{cccc}
Ar^{1} - A - \left[Ar^{2} - A - Ar^{3}\right] & Ar^{3}
\end{array}\right)$$

Formula 1

Wherein n is at least 1; each A is N or P; each Ar1 and Ar3 is arylene or heteroarylene; Ar2 is arylene or heteroarylene containing a linking ring to which the two A atoms are both directly linked and at least one of Ar1 or Ar2 is substituted with at least one substituent.

Antoniadis discloses formula A-1 (column 9, line 45):

$$A \longrightarrow Ar_1 \longrightarrow N \longrightarrow Ar_1 \longrightarrow A$$

$$Ar_2 \longrightarrow N \longrightarrow Ar_2 \longrightarrow Ar_2$$

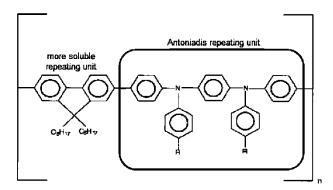
Formula A-1 shows n =1, N corresponds to applicants' A group, Ar1 corresponds to applicants' Ar1, Ar2 corresponds to applicants' Ar3 and the phenyl linking group corresponds to applicants' Ar2. The phenyl linking group connects both N atoms.

Formula A-1 also meets the limitations of claim 24.

Antoniadis fails to mention a second repeating unit wherein the second repeat unit is selected from optionally substituted phenyl, fluorene, spirobifluorene, indenofluorene, heteroaryl or dihydrophenanthrene.

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Bradley discloses that fluorene-triarylamine copolymers offer some advantages over a purely triarylamine based system. Bradley teaches that when N,N'-diphenyl, N,N'-bis(3-methylphenyl)-(1,1'-diphenyl)-4,4'-diamine) (TPD) is combined with a more soluble polymer (fluorene based) the resulting copolymer is more processible and has a higher hole mobility (page 242). The copolymers can be represented by the Formula B-1:



Formula B-1 disclosed by Bradley shows the same triarylamine based repeating unit as Antoniadis copolymerized with a fluorene derivative which is a more soluble repeating unit. Copolymer B-1 shows monomer 1 in conjugation with monomer 2 (per claim 7) and the second monomer is fluorenyl (per clam 27).

As Bradley teaches that such copolymer produce higher hole mobility than polymers with only the triarylamine component present, one could envisage adding a more soluble monomeric species to the polymers of Antoniadis to improve the processibility and promote higher hole mobility.

It would have been obvious to a person of ordinary skill in the art at the time of the invention to have modified the polymers of Antoniadis by adding the more soluble

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repeating units (fluorenyl) of Bradley since Bradley teaches that the resulting copolymer that read on the instant limitation improve processibility and hole mobility properties.

18. **Regarding Claims 13-14,** Antoniadis discloses Formula A-1 above where A can be a halogen atom (column 4, lines 21-22) which corresponds to applicants' LG group in Formula II (claim 13), n =1, N corresponds to applicants' A group, Ar1 corresponds to applicants' Ar1, Ar2 corresponds to applicants' Ar3 and the phenyl linking group corresponds to applicants' Ar2 linking both N atoms. The reactions can be catalyzed by palladium (variable oxidation state metal) (column 10, lines 29-30). The addition of a second repeating unit was discussed above.

Antoniadis also discloses that the reaction can be carried out in the presence of a catalytic amount of a divalent nickel salt (column 11, lines 23-26) (per claim 14).

- 19. **Regarding Claims 2-3 and 25-26**, Antoniadis discloses in Formula A-1 that the R substituent on the phenyl linking group (called Ar2 by applicant) can be a C1-24 hydrocarbyl (alkyl) group (abstract).
- 20. **Regarding Claim 4**, Antoniadis discloses that the phenyl linking group which corresponds to applicants' Ar2. Ar1 of Formula A-1 can be an aryl group C6-18 which includes a phenyl group (abstract).

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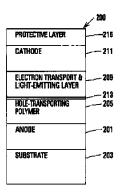
21. **Regarding Claims 5 and 9**, Antoniadis discloses that Ar2 in A-1 which corresponds to applicants' Ar3 can be substituted with C1-24 hydrocarbyl (alkyl) alkyl group (abstract).

- 22. **Regarding Claim 10**, Antoniadis discloses the polymers are combined with at least one organic compound and functions as a hole conducting (transporting) layer (abstract).
- 23. **Regarding Claims 15 and 16,** Antoniadis discloses that Formula A-1 (shown above) can form a copolymer with a second aromatic monomer as shown below (abstract):

The reactions can be catalyzed by palladium and sodium t-butoxide (base) (column 10, lines 29-30) with A (LG, leaving group) being chlorine or bromine (column 9, line 50 and column 8, lines 43-44). The A (LG, leaving group) can also be boronic acid derivatives catalyzed by palladium (column 13, lines 22-25).

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24. **Regarding Claims 17 and 19-20,** Antoniadis discloses that the conductive polymer composition can be used in an electroluminescent device (optical device) (abstract) (per claim 20). The hole conducting layer/hole transporting layer is formed from the poly (arylmine) (oligomer, per claim 17) (column 3, lines 65-66) (column 5, lines 1-3). The hole transporting layer is located between the anode and cathode(per claim19), see figure 1 below:



- 25. Claims 21 and 23 are rejected under 35 U.S.C. 103(a) as being unpatentable over Antoniadis (US 5,948,552) in view of Allen (US 6,858,703).
- 26. **Regarding Claim 21 and 23,** Antoniadis discloses that the copolymers can be used in an electroluminescent device (abstract) but fails to mention a switching device.

Allen discloses that triarylamine polymers are used in optical sensor, switching devices and field effect transistors (column 83, lines 20-30).

It would have been obvious to a person of ordinary skill in the art at the time of the invention to have used the electroluminescent device of Antoniadis in applications

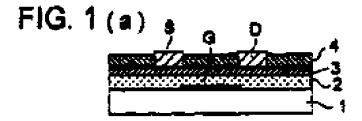
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taught by the prior art which would have included the applications disclosed by Allen which reads on the instant limitations.

- 27. Claim 22 is rejected under 35 U.S.C. 103(a) as being unpatentable over Antoniadis (US 5,948,552) in view of Allen (US 6,858,703) and further in view of Hirai (US 6,740,900).
- 28. **Regarding Claim 22,** Antoniadis in view of Allen disclose triarylamine polymers used as a field transistors but fail to mention the structure of the device.

Hirai discloses that an insulator layer can be placed in various locations with respect to the gate, drain and source electrodes in an organic thin-film transistor.

Hirai discloses that the organic thin-film transistor contains a organic semiconductor layer 3, a gate electrode G, a drain electrode D, and a source electrode S that are in Figure I(a) shown below:



Hirai further discloses that a dielectric layer (field-effect transistor) serves as gate insulation layer 2 (column 7, lines 44-45). The above figure shows that the gate electrode G and the organic semiconductor layer 3 are both in contract with the gate insulation layer 2 located on side one. In addition, the above figure shows that the source electrode S and the drain electrode D are located on the organic semiconductor layer 3 on side two.

Hirai shows in the prior art that the claimed structure can be use as a field effect transistor.

With the reasonable expectation of success a person with ordinary skill in the art at the time of the invention would have selected from known field-effect transistor structures and readily substituted the conductive polymers disclosed by Antoniadis (K-2) for the organic semiconductive layer 3 disclosed by Hirai in Figure I (a) since the conductive polymers of Antoniadis would be functional equivalents to the polymers typically used to make an organic semiconductive layer.

Response to Arguments

Applicant has amended some claims and added some new claims. Additionally, applicant argues that Antoniadis only teaches triarylamine amine polymer but not copolymers with the second monomers as listed in claims 1 and 13.

While Antoniadis does not mention a second monomer as claimed by applicant,

Bradley discloses that fluorene-triarylamine copolymers offer some advantages over a

purely triarylamine based system. Bradley teaches that when N,N'-diphenyl, N,N'-bis(3-methylphenyl)-(1,1'-diphenyl)-4,4'-diamine) (TPD) is combined with a more soluble polymer (fluorene based) the resulting copolymer is more processible and has a higher hole mobility (page 242). As Bradley teaches how such copolymer produce higher hole mobility than polymers with only the triarylamine component present, one could envisage adding a more soluble monomeric species to the polymers of Antoniadis to improve the processibility and promote higher hole mobility.

Additionally, the examiner has applied Towns who teaches triarylamine based copolymers that read on applicant monomer 1 and monomer 2 limitations.

The applicant's arguments with respect to claims 1-5, 7, 9-10, 13-27 have been considered but are most in view of the new grounds of rejection necessitated by the applicant's amendment.

Conclusion

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any

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extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

The prior art made of record and not relied upon is considered pertinent to applicant's disclosure. Wu (US 5,728,801).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to GREGORY CLARK whose telephone number is (571)270-7087. The examiner can normally be reached on M-Th 7:00 AM to 5 PM Alternating Fri 7:30 AM to 4 PM and Off.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Larry Tarazano can be reached on (571) 272-1515. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.